



RHODE ISLAND DEPARTMENT OF HEALTH  
DIVISION OF DISEASE PREVENTION AND CONTROL  
STD PROGRAM, Cannon Building, 3 Capital Hill, Room 106, PROVIDENCE, RI 02908  
TEL: (401) 222-2577 FAX: (401) 222-1105



**STD treatment information is very important. Because of this:**  
**If making copies of this form, please copy both sides. If faxing this form, please fax both sides.**

**CONFIDENTIAL REPORT FOR SEXUALLY TRANSMITTED DISEASES INSTRUCTIONS**

1. Mail or fax fully completed report within 4 days or as soon as treatment is prescribed.
2. FAX or Phone partial report immediately if partner services are requested (see VI below) or syphilis reporting criteria are met.

**I. PATIENT INFORMATION:**

Last Name		First (full name)		MI
Street			Apt. #	
City/Town		Zip Code	Phone Number & Area Code	
Sex <input type="checkbox"/> Male <input type="checkbox"/> Female	Date of Birth ____/____/____	Age	Marital Status <input type="checkbox"/> Single <input type="checkbox"/> Married <input type="checkbox"/> Other <input type="checkbox"/> Unknown	
Ethnic Origin: <input type="checkbox"/> Hispanic or Latino <input type="checkbox"/> Not Hispanic or Latino	Race (indicate one or more) <input type="checkbox"/> Black <input type="checkbox"/> White <input type="checkbox"/> Hawaiian Native <input type="checkbox"/> American Indian/Alaskan Native <input type="checkbox"/> Asian			

**II. FACILITY INFORMATION:**

Physician or Facility Name
Facility Contact Person for STD Reporting
Facility Street Address, Facility City, State, Zip
Facility Phone Number & Area Code

**III. RISK FACTORS:**

1. Sexual orientation of the patient: ☐ Heterosexual ☐ Homosexual ☐ Bisexual ☐ Unknown
2. How many sexual partners has the patient had in the past 6 months? \_\_\_\_\_
3. Has the patient participated in anonymous sex in the past 6 months? ☐ Yes ☐ No ☐ Unknown
4. How often had the patient used condoms in the past 6 months? ☐ Always ☐ Sometimes ☐ Never ☐ Unknown
5. Has the patient recruited sexual partners from high-risk locations (night clubs, parks, the internet...)? ☐ Yes ☐ No ☐ Unknown  
If yes, where? \_\_\_\_\_
6. Has the patient used Intravenous Drugs in the past 6 months? ☐ Yes ☐ No ☐ Unknown If yes, list: \_\_\_\_\_
7. Has the patient used Non-Intravenous Drugs in the past 6 months? ☐ Yes ☐ No ☐ Unknown If yes, list: \_\_\_\_\_
8. Has the patient supplied sex for drugs or money in the past 6 months? ☐ Yes ☐ No ☐ Unknown
9. Has the patient received sex for drugs or money in the past 6 months? ☐ Yes ☐ No ☐ Unknown
10. Is the patient co-infected with Hepatitis B or C? ☐ Yes ☐ No ☐ Unknown

**IV. PATIENT TREATMENT INFORMATION:**

Has the patient received client specific prevention counseling? <input type="checkbox"/> Yes <input type="checkbox"/> No	Is this patient pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No
Did this patient receive treatment? <input type="checkbox"/> Yes (If yes, check box on back for treatment administered) <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> Unable to reach	If yes, when did the patient receive treatment? Date ____/____/____

**V. STD INFORMATION:**

<b>1. GONORRHEA</b>		<b>2. CHLAMYDIA</b>	
Culture: Result____ Date____	<input type="checkbox"/> Cervical <input type="checkbox"/> Urethral <input type="checkbox"/> Rectal <input type="checkbox"/> Pharyngeal/Throat <input type="checkbox"/> Other_____	EIA: Result____ Date____	<input type="checkbox"/> Cervical <input type="checkbox"/> Urethral <input type="checkbox"/> Rectal <input type="checkbox"/> Other_____
Gram Stain: Result____ Date____		Urine DNA: Result____ Date____	
Urine DNA: Result____ Date____			Did the patient have PID <input type="checkbox"/> Yes <input type="checkbox"/> No
LCR (or equivalent) Result____ Date____	Did the patient have PID? <input type="checkbox"/> Yes <input type="checkbox"/> No		
Other Test _____			
<b>3. SYPHILIS</b>		<b>4. PID</b>	
RPR Titer _____ Date____	<input type="checkbox"/> Primary (Lesion) <input type="checkbox"/> Secondary (Rash, Other Symptoms) <input type="checkbox"/> Early Latent (Asymptomatic, less than 1 year) <input type="checkbox"/> Late Latent (over 1 year duration) <input type="checkbox"/> Neurosyphilis <input type="checkbox"/> Congenital (infant)	<input type="checkbox"/> In-Patient <input type="checkbox"/> Out-Patient	<input type="checkbox"/> Gonococcal <input type="checkbox"/> Chlamydia <input type="checkbox"/> Agent Unknown <input type="checkbox"/> Other: _____
FTA Result _____ Date____			

**5. OTHER STDs**

- ☐ Chancroid ☐ Granuloma Inguinale ☐ Lymphogranuloma- Venereum (LGV)

**VI. PARTNER NOTIFICATION:**

Providers treating STD's are expected to counsel patients in prevention and identify and refer partners to medical care for examination and treatment.

- Was the patient given partner notification information? ☐ Yes ☐ No
  - When partners are hard to reach, the Department of Health STD program can provide this confidential service on a very limited basis.
  - Are you requesting state resources for partner services for this case? ☐ Yes ☐ No
- Fax immediately and we will call you to get more details.

## SUMMARY OF THE 2002 CDC SEXUALLY TRANSMITTED DISEASES (STD) TREATMENT GUIDELINES STD CONTROL PROGRAM – RHODE ISLAND DEPARTMENT OF HEALTH

These guidelines for the treatment of STDs reflect the recommendations of the 2002 CDC STD Treatment Guidelines. These are outlines for quick reference that focus on STDs encountered in an outpatient setting and are not an exhaustive list of effective treatments. Please refer to the complete document of the CDC for more information or call the STD Program. These guidelines are to be used for clinical guidance and are not to be construed as standards or inflexible rules. Clinical and epidemiological services are available through the STD Program and staff is also available to assist healthcare providers with confidential notification of sexual partners of patients infected with STDs and HIV. Please call for any assistance at: (401) 222-2577. FAX (401) 222-1105. STD Program, Rhode Island Department of Health, Cannon Building, 3 Capital Hill, Room 106, Providence, RI 02908

Disease	Recommended Treatment	Alternative
<b>SYPHILIS (see CDC guidelines for follow-up recommendations)</b>		
Primary, Secondary or Early Latent (<1 Year) Adults  Children	<input type="checkbox"/> Benzathine penicillin G 2.4 million units IM in a single dose  <input type="checkbox"/> Benzathine penicillin G 50,000 units/kg IM, up to the adult dose of 2.4 million units in a single dose	<b>(For penicillin allergic, non-pregnant <u>adult</u> patients)</b> <input type="checkbox"/> Doxycycline 100 mg orally 2 times a day for 14 days <b>OR</b> <input type="checkbox"/> Ceftriaxone 1 g daily IV or IM for 8-10 days <b>OR</b> <input type="checkbox"/> Azithromycin 2 g orally single dose
Late Latent (>1 Year) or Latent of Unknown Duration  Adults  Children	<input type="checkbox"/> Benzathine penicillin G 2.4 million units IM <b>for 3 doses</b> , 1 week apart (total 7.2 million units)  <input type="checkbox"/> Benzathine penicillin G 50,000 units/kg IM up to the adult dose of 2.4 million units, administered as three doses at 1 week intervals (total 150,000 units up to the adult total dose of 7.2 million units)	<input type="checkbox"/> Doxycycline 100 mg orally 2 times a day for 28 days <b>for adults only</b>
Neurosyphilis	<input type="checkbox"/> Aqueous crystalline penicillin G 18- 24 million units per day, administered as 3-4 million units IV every 4 hours or continuous infusion, for 10-14 days	<input type="checkbox"/> Procaine penicillin 2.4 million units IM once daily <b>plus</b> probenecid 500 mg orally 4 times a day, both for 10-14 days
HIV Infection	<b>For primary, 2<sup>nd</sup> and early latent syphilis:</b> Treat as above. Some specialists recommend three doses. <b>For late latent syphilis or syphilis of unknown duration:</b> perform CSF examination before treatment	
Pregnancy	<b>Penicillin is the <u>only</u> recommended treatment for syphilis during pregnancy. Women who are allergic should be desensitized and then treated with penicillin. Dosages are the same as in non-pregnant patients for each stage of syphilis.<sup>1</sup></b>	
<b>GONOCOCCAL INFECTIONS<sup>2</sup></b>		
Cervix, Urethra, Rectum   Pharynx  Conjunctiva	<input type="checkbox"/> Ceftriaxone 125 mg IM once <b>OR</b> <input type="checkbox"/> Cefixime 400 mg orally once <b>OR</b> <input type="checkbox"/> Ciprofloxacin <sup>4,5</sup> 500 mg orally once <b>OR</b> <input type="checkbox"/> Ofloxacin <sup>4,5</sup> 400 mg orally once <b>OR</b> <input type="checkbox"/> Levofloxacin <sup>4,5</sup> 250 mg orally once  <input type="checkbox"/> Ceftriaxone 125 mg IM once <b>OR</b> <input type="checkbox"/> Ciprofloxacin <sup>4,5</sup> 500 mg orally once  <input type="checkbox"/> Ceftriaxone 1 g IM once plus lavage the infected eye with saline solution once	<input type="checkbox"/> Spectinomycin <sup>3</sup> 2 g IM once (see CDC guidelines for other cephalosporins and quinolones)
<b>Children (&lt;45KG)</b> Vagina, Cervix, Urethra, Pharynx, Rectum	<input type="checkbox"/> Ceftriaxone 125 mg IM once	<input type="checkbox"/> Spectinomycin <sup>3</sup> 40 mg/kg IM once (maximum 2 g)
<b>Neonates</b> Ophthalmia Neonatorum <sup>6</sup> Infants born to infected mothers	<input type="checkbox"/> Ceftriaxone 25-50 mg/kg IV or IM once (maximum 125 mg)	
<b>Pregnancy</b>	<input type="checkbox"/> Ceftriaxone 125 mg IM once	<input type="checkbox"/> Spectinomycin <sup>3</sup> 2 g IM once
<b>CHLAMYDIAL INFECTIONS</b>		
<b>Adult</b>	<input type="checkbox"/> Azithromycin 1 g orally single dose <b>OR</b> <input type="checkbox"/> Doxycycline 100 mg orally 2 times a day for 7 days	<input type="checkbox"/> Erythromycin base 500 mg orally 4 times a day for 7 days <b>OR</b> <input type="checkbox"/> Erythromycin ethylsuccinate 800 mg orally 4 times a day for 7 days <b>OR</b> <input type="checkbox"/> Ofloxacin <sup>4</sup> 300 mg orally 2 times a day for 7 days <b>OR</b> <input type="checkbox"/> Levofloxacin <sup>4</sup> 500 mg orally once a day for 7 days
<b>Children</b> ≤ 45 kg----->  ≥ 45 kg and < 8 Years of Age-----> ≥ 8 Years of Age----->	<input type="checkbox"/> Erythromycin base or ethylsuccinate 50 mg/kg/day orally divided into four doses daily for 14 days <sup>7</sup> <input type="checkbox"/> Azithromycin 1 g orally single dose <input type="checkbox"/> Azithromycin 1 g orally single dose <b>OR</b> <input type="checkbox"/> Doxycycline 100 mg orally 2 times a day for 7 days	
<b>Pregnancy</b>	<input type="checkbox"/> Erythromycin base 500 mg orally 4 times a day for 7 days <b>OR</b> <input type="checkbox"/> Amoxicillin 500 mg orally 3 times a day for 7 days	<input type="checkbox"/> Erythromycin 250 mg orally 4 times a day for 14 days <b>OR</b> <input type="checkbox"/> Erythromycin ethylsuccinate 800 mg orally 4 times a day for 7 days (or 400 mg 4 times a day for 14 days) <b>OR</b> <input type="checkbox"/> Azithromycin 1 g orally single dose

<sup>1</sup> Tetracycline/doxycycline contraindicated; erythromycin not recommended because it does not reliably cure an infected fetus; data insufficient to recommend azithromycin or ceftriaxone.

<sup>2</sup> Treat also for *Chlamydia trachomatis* if not ruled out by a sensitive test.

<sup>3</sup> Not effective against incubating syphilis and is less effective against pharyngeal gonorrhea.

<sup>4</sup> Quinolones are contraindicated in pregnant women. No joint damage attributable to quinolone therapy has been observed in children treated with prolonged ciprofloxacin regimens. Thus children who weigh ≥ 45 kg can be treated with any regimen recommended for adults.

<sup>5</sup> Quinolones should not be used for gonococcal infections acquired in Asia or the Pacific, including Hawaii. In addition, use of quinolones is probably inadvisable for treating infections acquired in California and in other areas with increased prevalence of quinolone resistance.

<sup>6</sup> Hospitalize and evaluate disseminated infection.

<sup>7</sup> The efficacy of treating neonatal chlamydia conjunctivitis and pneumonia is about 80%. A second course of therapy may be required. An association between oral erythromycin and infantile hypertrophic pyloric stenosis has been reported in infants less than 6 weeks treated with this drug. See CDC guidelines for more information.